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The attached documents are exact copies of the European patent application conformes à la version described on the following page, as originally filed.

Les documents fixés à cette attestation sont initialement déposée de la demande de brevet européen spécifiée à la page suivante.

Patentanmeldung Nr.

Patent application No. Demande de brevet n°

99200353.3

Der Präsident des Europäischen Patentamts; Im Auftrag

For the President of the European Patent Office

Le Président de l'Office européen des brevets p.o.

W. DRAISMA

DEN HAAG, DEN THE HAGUE, LA HAYE, LE

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Blatt 2 der Bescheinigung Sheet 2 of the certificate Page 2 de l'attestation

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Centrifugal atomized zinc alloy powders in alkaline batteries

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CENTRIFUGAL ATOMIZED ZINC ALLOY POWDERS IN ALKALINE BATTERIES

This invention relates to centrifugal atomized zinc alloy powders for alkaline batteries.

Zinc alloy powders are used as an ingredient of the anode active material of alkaline batteries. Zinc alloy powders containing one or more elements selected from In, Bi, Al, Ca and Pb, are known from US 5,082,622 and

10 WO 94/19502. Such alloys are normally prepared by traditional powder manufacturing techniques, such as allowing the molten alloy to flow away in a jet of compressed air. The use of a protective atmosphere in such gas jet atomisation process is possible, but quite expensive because of the high flow rate needed. Powders which are manufactured by compressed air atomisation do not always have an acceptable resistance to corrosion in the electrolyte of the battery, especially after partial discharge of the latter. A well-known way of dealing with a low resistance to corrosion is by adding certain amounts of organic or inorganic products. An example of an organic product is polyethylene glycol. However this procedure also has some disadvantages, e.g. the battery capacity could be decreased.

Centrifugal atomisation is a method in which molten metal is poured from a nozzle on a high velocity rotating plate, a cup or a disk with a predetermined shape whereby the metal is disintegrated by centrifugal force. The above cited documents merely mention that the powders can also be manufactured by centrifugal atomisation without specifying any working condition. In these documents the particular effects of using centrifugal atomized powders are not investigated.

According to the invention, a zinc alloy powder prepared by centrifugal atomisation in a protective atmosphere with a reduced oxygen content has a better resistance to corrosion than a powder prepared by atomisation in a stream of air. Moreover, unacceptably low corrosion resistance values of some of the above mentioned zinc alloy powders prepared by atomisation in air are turned into acceptable values when the same alloys are prepared by centrifugal atomisation in said protective atmosphere. The use of organic and inorganic corrosion inhibitors can be limited, or even eliminated. The capacity of batteries containing powders according to the invention is good. The production method

10 and Ca, or

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is economically sound, because the consumption of gas in the protective atmosphere is very low.

The alloy powder of the invention consists of either

(a) 0.005-2 %, and preferably 0.01-2 % by weight of indium; and 0.005-0.2 %, preferably 0.01-0.2 % by weight of either one of Al and Bi, or (b) 0.005-2 % by weight of indium; and 0.005-0.2%, preferably 0.01-0.2 % by weight of Bi; and 0.001-0.5%, preferably 0.003-0.5 % of either one or both of Al

(c) 0.005-2%, and preferably 0.01-2 % by weight of either one or both of Bi and Al,

and in each case 0-0.5 % by weight of Pb, the remainder being zinc.

15 By zinc is meant here and in the following thermally or electrolytically refined zinc (Special High Grade). For economical reasons the indium content can be limited to 5000 ppm, or even to 1000 ppm.

The powder is prepared by centrifugal atomisation, characterized in that this atomising process is performed in a protective atmosphere, mainly consisting of inert gasses, which controls the oxidation. More particularly, the oxygen content in the atmosphere is less than 4% by.

In order to prevent excessive oxidation, the oxygen content is limited to 4%, and preferably less than or equal to 3.5%. On the other hand, a minimum amount of oxygen can be useful to control the shape of the solidified droplets.

25 Also, too low oxygen contents imply a reduction of the capacity of the battery. For these reasons the oxygen content is preferably greater than 0%, and especially greater than or equal to 0.2%.

The rotating disk of the atomizer is made out of materials like alumina, carbon or fused silica, which can also be coated to prevent oxidation or occurrence of accretion.

The applicant has found that when the powders described previously are prepared by a centrifugal atomising process in the above described protective atmosphere, they are consistently showing a better resistance to corrosion in the electrolyte of the battery, after partial discharge of the latter, compared to powders with identical compositions and prepared by gas jet atomisation. The resistance to corrosion and the capacity of the powders according to the

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compositions of the invention, and prepared by centrifugal atomisation, are all recommendable for practical use in alkaline batteries.

In case the desired powder contains additives which can be cemented (for instance In), then another way to produce the powder consists in preparing a powder with the additives which cannot be cemented and optionally with a part of the additives which can be cemented according to the above mentioned method of centrifugal atomisation and making an anode from the thus obtained powder. That anode is introduced in the battery and the additives which can be cemented, are either added to the electrolyte of the battery, from which they 10 cement on the powder of the anode, or they are cemented before the introduction in the battery. It is also possible to obtain powders for alkaline batteries by mixing a powder with a given composition and produced in a traditional way, such as atomisation in a stream of air, with a powder which was prepared by centrifugal atomisation in a protective atmosphere, according to the invention.

This invention relates not only to a powder such as it is introduced in the battery, but also to a powder such as it is present in the battery, and which consists entirely or partly of powder prepared by centrifugal atomisation, the remainder being prepared by traditional powder manufacturing techniques.

The powder is suitable for use in any type of battery, such as LR6, LR14, LR20.

Here, the following should be noted: Centrifugal atomized zinc alloy powders are mentioned in "Atomization of melts for Powder Production and Spray 25 Deposition", by A.J. Yule and J. J. Dunkley; Clarendon Press, Oxford; 1994; pp. 223-224. Centrifugal atomisation is used as it gives a very closely controlled particle size distribution, which renders the powder production more economical. It is also possible, by control of the atmosphere inside the atomizer, to control the particle shape and hence the surface area and apparent 30 density of the powder, both of which are significant for battery applications. This publication however does not mention which alloys can be exploited beneficially for this application, and neither is described what the controlled atmosphere inside the atomizer should be. Also it does not describe if and in what way the process of centrifugal atomisation has an influence on the characteristics of the batteries prepared with these powders.



In an example of producing the powder of the invention, the powders are prepared by adding the desired quantities of the alloying elements to molten zinc, which is homogenized by stirring at elevated temperature. The process of centrifugal atomisation is carried out as follows: about 150 kg of alloyed zinc is melted in a protective atmosphere at 480°C and poured on a rotating disk. The rotating disk is made of carbon, and placed 25 mm below the ejecting nozzle concentrically, 1.5 m above the bottom at the centre of the atomising chamber. The atomising chamber is filled with nitrogen and 0.4% of oxygen. The rotation speed is about 5000 rpm, the diameter of the disk is 170 mm and the flow rate of 10 molten-metal-is-about 530 kg/hr. The produced powders are collected and their particle size distribution is verified.

With the alloy powders batteries of the LR14-type are produced. These batteries are discharged over 2 Ohm for 1 hr or for 4 1/2 hrs. Subsequently one determines the quantity of hydrogen which is evolved when the batteries are maintained for 7 days at 71°C. The resulting gas production is expressed as ml of gas per battery. Also the volume expansion of a partially discharged battery is measured after 24 hrs at 71°C, and the volume increase is expressed as a percentage of the original volume. This measuring technique is described in US 5,364,715.

The results of these tests are compared with results on air atomized powders with the same chemical composition. In the examples, all powders are sieved on 500 micron to take away the very course particles.



The results are summarized in the following table:

No.	Alloy powder additives (ppm)				Prepara- tion (*)	Volume expan- sion (%)	Gassing after partial discharge 4 1/2 hrs, 2 Ohm (ml)	Gassing after partial discharge 1 hr, 2 Ohm (ml)
	In	Bi	Ca	Al				
(1)	150	230	-	-	С	3.4	4.63	1.95
(2)	500	100	-	100	С	1.7	2.07	1.5
(3)	200	180	-	70	С	3.6	3.67	1.7
(4)	250	250	150	-	¢	3.1	3.57	1.37
(5)	-	250	-	-	С	6.5	8.05	3.12
(6)	2000	500	-	-	C	3.4	2.38	1.83
(7)	150	230	-	-	T	28.4	10.65	13.63
(8)	500	100	-	100	T	5.6	2.5	2.3
(9)	200	180	-	70	T	5.1	4	1.7
(10)	250	250	150	- 3	T	10.2	4.08	6.75
(11)	-	250	-	- 3	T	19.8	9.23	12.13
(12)	2000	500	-	- 1	T	41.9	12.7	19.58
(13)	500	5000	-	<i>7</i> 0	T	26.2	16.7	13.5
(14)	500	5000	-	70	С	30.1	13.5	6.37
(15)	600		-	-	T	87.6	52.63	(**)
(16)	600	-	-	-	С	42.1	25.07	22.43
(17)	900	-	-	- //	С	25.9	21.38	15.9
(18)	40	40		-	С	28.5	11.63	8.57

(*): 'C': centrifugal atomized powder; 'T':powder preparation by gas jet atomisation

(**): Leakage occurred, which indicates that gassing was excessive and unacceptable.

For the volume expansion, a value of less than 12% is considered to be very good, preferably the expansion should be less than 8%.

The gassing after partial discharge is good for values less than 12 ml, and preferably less than 8 ml, for both the gassing tests after 4½ hrs or 1 hr of discharge.

Comparison of examples No. (1)-(6) with examples No. (7)-(12) shows that the powders according to the invention have better corrosion resistance and expansion parameters than traditionally prepared powders. The compositions of the examples No. (7), (11) and (12), which do not yield acceptable results when prepared in the traditional way, have excellent results when prepared according to the invention, as shown in examples No. (1), (5) and (6). Examples No. (14), (16)-(18) show that centrifugal atomisation is not yielding satisfactory results for zinc powders consisting of alloys other than the compositions claimed in the invention.

10 It has been noticed that the powder capacity is varying between 90 mand-110 % as compared to a standard powder used in LR6 batteries nowadays, for instance the powder according to example No. (9) above.



CLAIMS

- A centrifugal atomized zinc alloy powder for alkaline batteries consisting of either of
- 5 (a) 0.005-2 % by weight of indium, and 0.005-0.2 % by weight of either one of Al and Bi, or
 - (b) 0.005-2 % by weight of indium, and 0.005-0.2 % by weight of Bi, and 0.001-0.5 % of either one or both of Al and Ca, or
 - (c) 0.005-2 % by weight of either one or both of Bi and Al,
- and 0-0.5 % by weight of Pb, the remainder being zinc, and characterized in that the centrifugal atomising process is performed in a protective atmosphere, where the oxygen content is less than 4% by volume.
- 2. A centrifugal atomized zinc alloy powder according to claim 1, consisting of
 (a) 0.01-2 % by weight of indium, and 0.01-0.2 % by weight of either one of Al and Bi, or
 - (b) 0.005-2 % by weight of indium, and 0.01-0.2 % by weight of Bi, and 0.003-0.5 % of either one or both of Al and Ca, or
 - (c) 0.01-2 % by weight of either one or both of Bi and Al, and 0-0.5 % by weight of Pb, the remainder being zinc.
 - 3. A centrifugal atomized zinc alloy powder in alkaline batteries according to claim 1 or 2, characterized in that the oxygen content in the protective atmosphere is greater than 0 % by volume.
- 4. A centrifugal atomized zinc alloy powder in alkaline batteries according to
 claim 3, characterized in that the oxygen content in the protective atmosphere is between 0.2 % and 3.5% by volume.
 - 5. An alkaline battery consisting of an anode, a cathode and an electrolyte, characterized in that the battery uses a centrifugal atomized zinc alloy powder according to one of the claims 1-4.
- 30 6. An alkaline battery according to claim 5, characterized in that the powder comprises metal cemented out of the electrolyte.

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ABSTRACT

USE OF CENTRIFUGAL ATOMIZED ZINC ALLOY POWDERS IN ALKALINE BATTERIES

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This invention relates to centrifugal atomized zinc alloy powders for alkaline batteries consisting of

- (a) 0.005-2 % by weight of indium, and 0.005-0.2 % by weight of either one of Al and Bi, or
- (b) 0.005-2 % by weight of indium, and 0.005-0.2 % by weight of Bi, and 0.001-0.5 % of either one or both of Al and Ca, or
 (c) 0.005-2 % by weight of either one or both of Bi and Al, and 0-0.5 % by weight of Pb, the remainder being zinc.
- 15 The powder is obtained by centrifugal atomisation in a protective atmosphere, where the oxygen content is less than 4% by volume.
- The resistance to corrosion in the electrolyte of the battery, especially after partial discharge, is markedly better than when the same alloys are prepared by the traditional production process. The capacity of batteries containing these powders is very good.

